Please replace formula IIIaf on page 28 with the following rewritten formula:

Please replace formula IVo on page 32 with the following rewritten formula:

Please replace formula Vi on page 34 with the following rewritten formula:

Please replace formula Vt on page 36 with the following rewritten formula:

IN THE CLAIMS:

Please amend claims 11, 13, 14, 33, 34, 36, and 38, and enter new claims 41 and 42, as follows:

Sub-

11. (Amended) A method for providing sedation, mitigating anxiety or providing anaesthesia in a subject in need thereof, comprising administering to a subject an effective amount of a therapeutic compound, wherein the therapeutic compound is of the formula (Ia):

(Ia) $\begin{array}{c}
G^{2} \\
| \\
G^{1}
\\
| \\
F^{-}CF^{2}
\\
| \\
FONO_{2}
\end{array}$

NDI

in which F^2 is an organic radical which may be joined in a cyclic ring system with G^2 , and which may contain inorganic counterions, but is not a nitrate group; E is a methylene group and G^1 is a methylene group or does not exist; F^1 is H; and G^2 is $R^N Z^N$;

wherein R^N is an organic radical possessing a heteroaryl group containing P or S atoms where said P or S are positioned β , γ , or δ to a nitrate group as identified in formula I; and Z^N is W^N_{mm}-X^N_m-Y^N_{oo};

wherein mm, nn, oo are 0 or 1 and WN, XN, YN are NH, NRNN, CO, O or CH2; wherein RNN is a short chain alkyl group $(C_1 - C_{12})$.

13. (Twice Amended) A method for providing sedation, mitigating anxiety or providing anaesthesia in a subject in need thereof, comprising administering to a subject an effective amount of a therapeutic compound, wherein the therapeutic compound is of the formula (Ic):

122

in which E is $(R^1R^2C)_m$ and $G^2-G^1-CF^1F^2-$ is $R^{19}-(R^3R^4C)_p-(R^{17}R^{18}C)_{n-}$;

wherein:

m, n, p are integers from 0 to 10;

R^{3,17} are each independently hydrogen, a nitrate group, or A; and R^{1,4} are each independently hydrogen, or A;

(122 Un.

where A is selected from a substituted or unsubstituted aliphatic group (comprising a branched or straight-chain aliphatic moiety having from 1 to 24/carbon atoms in the chain, which optionally may contain O, S, NR6 and unsaturations in the chain, optionally bearing from 1 to 4 hydroxy, nitrate, amino, aryl, or heterocyclic groups; an unsubstituted or substituted cyclic aliphatic moiety having from 3 to 7 carbon atoms in the aliphatic ring, which optionally may contain O, S, NR6 and unsaturations in the ring, optionally bearing from 1 to 4 hydroxy, nitrate, amino, aryl, or heterocyclic groups; an unsubstituted or substituted aliphatic moiety constituting a linkage of from 0 to 5 carbons, between R1 and R3 and/or between R17 and R4, which optionally may contain O, S, NR6 and unsaturations in the linkage, and optionally/bearing from 1 to 4 hydroxy, nitrate, amino, aryl, or heterocyclic groups); a substituted or unsubstituted aliphatic group (comprising a branched, cyclic or straight-chain aliphatic moiety having from 1 to 24 carbon atoms in the chain) containing carbonyl linkages (C=O, C=S, C=NOH), which optionally may contain O, S, NR6 and unsaturations in the chain, optionally bearing from 1 to 4 hydroxy, nitrate, amino, aryl, or heterocyclic groups; a substituted or unsubstituted aryl group; a heterocyclic group; an amino group selected from alkylamino, dialkylamino, cyclic amino, diamino and triamino moieties, arylamino, diarylamino, and alkylarylamino; hydroxy,/alkoxy; a substituted or unsubstituted aryloxy;

wherein X is F, Br, Cl, NO₂, CH₂, CF₂, O, NH, NMe, CN, NHOH, N₂H₃, N₂H₂R¹³, N₂H₃R¹⁴, N₃, S, SCN, SCN₂H₄(R¹⁵)₂/SCN₂H₃(R¹⁵), SC(O)N(R¹⁵)₂, SC(O)NHR¹⁵, SO₃M, SH, SR⁷, SO₂M, S(O)R⁸, S(O)₂R⁹, S(O)OR⁸, S(O)₂OR⁹, PO₂HM, PO₃HM, PO₃M₂, P(O)(OR¹⁵)(OR¹⁶), P(O)(OR¹⁶)(OM), P(O)(R¹⁵)(OR⁸), P(O)(OM)R¹⁵, CO₂M, CO₂H, CO₂R¹¹, C(O), C(O)R¹², C(O)(OR¹³), PO₂H, PO₂M, P(O)(ØR¹⁴), P(O)(R¹³), SO, SO₂, C(O)(SR¹³), SR⁵, SSR⁷ or SSR⁵;

Y is F, Br, Cl, CH₃, CF₂H, CF₃, OH, NH₂, NHR⁶, NR⁶R', CN, NHOH, N₂H₃, N₂H₂R¹³, N₂H₂R¹³, N₂H₃R¹⁴, N₃, S, SCN, SCN₂H₂(R¹⁵)₂, SCN₂H₃(R¹⁵), SC(O)N(R¹⁵)₂, SC(O)NHR¹⁵, SO₃M, SH, SR', SO₂M, S(O)₂R⁹, S(O)₂R⁹, S(O)₂OR⁹, PO₂HM, PO₃M₂, P(O)(OR¹⁵)(OR¹⁶), P(O)(OR¹⁶)(OM), P(O)(R¹⁵)(OR⁶), P(O)(OM)R¹⁵, CO₂M, CO₂H, CO₂R¹¹, C(O)R¹², C(O)(OR¹³), C(O)(SR¹³), SR⁵, SSR⁷ or SSR⁵, or does not exist;

 R^2 , R^5 , R^{18} , R^{19} are optionally hydrogen, A or X-Y;

R⁶, R⁷, R⁸, R⁹, R¹¹, R¹², R¹³, R¹⁴, R¹⁵, R¹⁶ are the same or different alkyl or acyl groups containing 1-24 carbon atoms which may contain 1-4 ONO₂ substituents; or C₁ - C₆ connections to R¹ - R⁴ in cyclic derivatives which may contain 1-4 ONO₂ substituents; or are each independently hydrogen, a nitrate group or A;

C 22 Person

M is H, Na+, K+, NH4+, N+H_kR¹¹(4_k) where k is 0-3; or other pharmaceutically acceptable counterion;

and with the proviso that when n = p = 1 and R^{19} , R^2 , R^{18} , $R^1 = H$ and R^{17} , R^3 are nitrate groups, R^4 is not H.

14. (Amended) The method of claim 11, wherein F² is a nitrate group; and E, F¹, G¹, G² are the same or different organic radicals which may be joined in cyclic ring systems, and which may contain inorganic counterions;

with the proviso that when E and G^1 are members groups and F^1 is H, G^2 is not a nitrate group, nor R^NZ^N ;

wherein R^N is any aryl or heteroaryl group and Z^N is (CO)_{mm}-X^N_{nn}-Y^N_{oo}; wherein mm, nn, oo are 0 or 1 and XN,Y^N are NH, NR^{NN}, O or CH₂; wherein R^{NN} is a short chain alkyl group ($C_1 - C_{12}$).

33. (Amended) A method of providing sedation or mitigating anxiety in a subject in need thereof, comprising administering to a subject an effective amount of a therapeutic compound selected from the group consisting of:

p 24

IIIb

IIIc

IIIe

Шf

IIIg

154

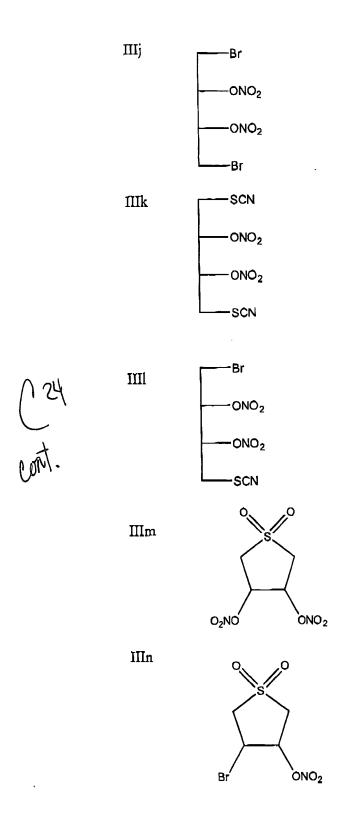
cont.

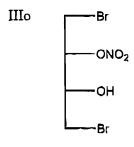
IIIh

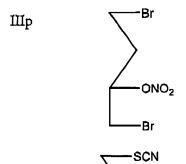
$$O_2NO$$
 O_2
 O_2NO
 O_2
 O_2
 O_2
 O_2
 O_2
 O_2
 O_2

Щі

9





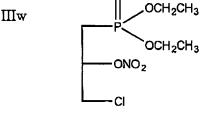


IIIr
$$S_2O_3$$
Na $-ONO_2$ ONO_2 S_2O_3 Na

,

IIIu
$$ONO_2$$
 SO_2H

Con-



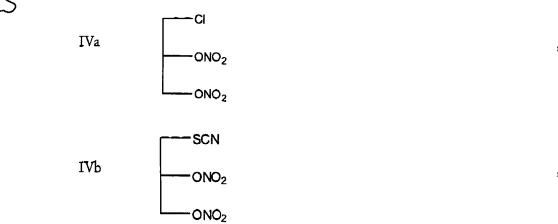
$$_{\mathrm{IIIz}}$$
 $_{\mathrm{O_{2}NO}}$ $_{\mathrm{NO_{3}H}}$

IIIah
$$O_2N$$
 O_2NO_2

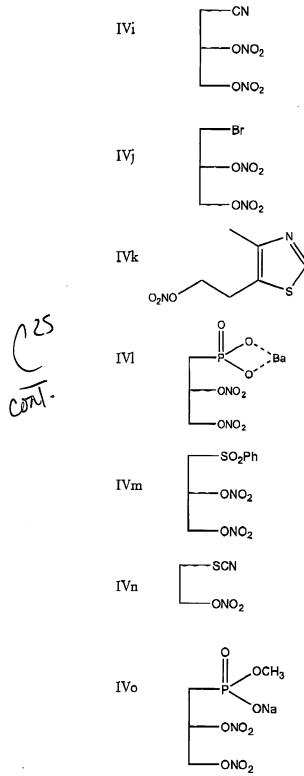
IIIai
$$O_2NO$$
 S_2O_3Na ONO_2

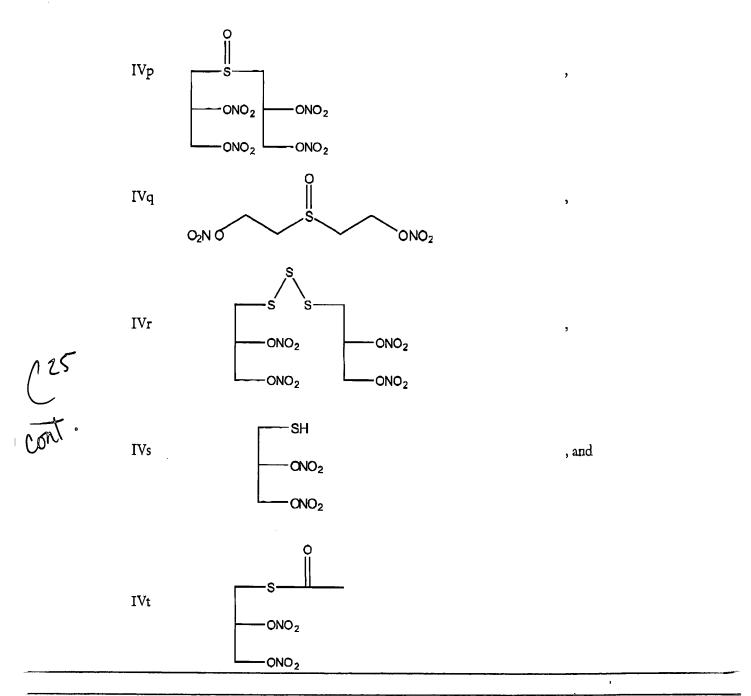
34. (Amended) The method of claim 33, wherein the compound has the formula IIIt:

36. (Amended) A method of providing sedation or mitigating anxiety in a subject in need thereof, comprising administering to a subject an effective amount of a therapeutic compound selected from the group consisting of:



$$S_2O_3Na$$
 ONO_2
 ONO_2





38. (Amended) A method of mitigating anxiety in a subject in need thereof, comprising administering to a subject an effective amount of a therapeutic compound selected from the group consisting of:

Va S—S—ONO₂ ONO₂ ONO₂

<u>∩</u> 26

Vb
$$O_2NO$$
 ONO_2

Vc ONO_2

Vd ONO_2

Ve ONO_2

Ve ONO_2

Ve S—S—CH₃

ONO₂

ONO₂

Vf S—S—OCH₃
ONO₂

Vg S—S—CI
ONO₂
ONO₂

Vh $S \longrightarrow S \longrightarrow NO_2$ ONO_2 ONO_2 ONO_2 ONO_2

,

Vj

 $V\mathbf{k}$

Cont.

Vm

Vn

